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IS 15938 (2011): Meat and Products -- Detection of Colouring Agents -- Method Using Thin-Layer Chromatography [FAD 18: Slaughter House and Meat Industry]



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IS 15938 : 2011
ISO 13496 : 2000

भारतीय मानक

मांस एवं मांस उत्पाद — रंजन अभिकर्मक का संसूचन — तनु-परत
क्रोमेटोग्राफी पद्धति द्वारा

Indian Standard

MEAT AND MEAT PRODUCTS — DETECTION OF
COLOURING AGENTS — METHOD USING
THIN-LAYER CHROMATOGRAPHY

ICS 67.120.10

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NEW DELHI 110002

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NATIONAL FOREWORD

This Indian Standard which is identical with ISO 13496 : 2000 'Meat and meat products — Detection of colouring agents — Method using thin-layer chromatography' issued by the International Organization for Standardization (ISO) was adopted by the Bureau of Indian Standards on the recommendation of the Slaughter House and Meat Industry Sectional Committee and approval of the Food and Agriculture Division Council.

The text of ISO Standard has been approved as suitable for publication as an Indian Standard without deviations. Certain conventions are, however, not identical to those used in Indian Standards. Attention is particularly drawn to the following:

- a) Wherever the words 'International Standard' appear referring to this standard, they should be read as 'Indian Standard'.
- b) Comma (,) has been used as a decimal marker while in Indian Standards, the current practice is to use a point (.) as the decimal marker.

The technical committee has reviewed the provisions of the following International Standards referred in this adopted standard and has decided that they are acceptable for use in conjunction with this standard:

<i>International/Other Standard</i>	<i>Title</i>
ISO 3696 : 1987	Water for analytical laboratory use — Specification and test methods
AOAC 46.1.08 : 1995	Official Methods of Analysis (AOAC International)

For the purpose of deciding whether a particular requirement of this standard is complied with, the final value, observed or calculated, expressing the result of a test or analysis, shall be rounded off in accordance with IS 2 : 1960 'Rules for rounding off numerical values (*revised*)'. The number of significant places retained in the rounded off value should be the same as that of the specified value in this standard.

Indian Standard

MEAT AND MEAT PRODUCTS — DETECTION OF COLOURING AGENTS — METHOD USING THIN-LAYER CHROMATOGRAPHY

1 Scope

This International Standard specifies a thin-layer chromatographic method for the detection of synthetic, water-soluble colouring agents in meat and meat products.

The following colouring agents can be detected with the method:

Tartrazine	Patent Blue V
Quinoline Yellow	Indigotine
Sunset Yellow FCF	Brilliant Black PN
Amaranth	Black 7984
Ponceau 4R	Fast Green FCF
Erythrosine	Blue VRS

Synonyms and identity numbers of these colouring agents are listed in annex A.

The plant colours and plant extracts which have been observed not to interfere with this method are listed in Table B.1. Natural colours which in some cases have been shown to interfere with this method are listed in Table B.2.

2 Normative references

The following normative documents contain provisions which, through reference in this text, constitute provisions of this International Standard. For dated references, subsequent amendments to, or revisions of, any of these publications do not apply. However, parties to agreements based on this International Standard are encouraged to investigate the possibility of applying the most recent editions of the normative documents indicated below. For undated references, the latest edition of the normative document referred to applies. Members of ISO and IEC maintain registers of currently valid International Standards.

ISO 3696:1987, *Water for analytical laboratory use — Specification and test methods*.

AOAC 46.1.08:1995, *Official Methods of Analysis* (AOAC International).

3 Term and definition

For the purposes of this International Standard, the following term and definition apply.

3.1

detection of colouring agents

detection of the presence or absence of colouring agents in accordance with the method specified in this International Standard

4 Principle

The colouring agents are extracted from a test portion with hot water and adsorbed onto polyamide powder. The extracted colouring agents are purified by column chromatography and the colours are eluted from the column. The colouring agents are identified by thin-layer chromatography.

5 Reagents

Use only reagents of recognized analytical grade, unless otherwise specified.

5.1 Water, complying with at least grade 3 in accordance with ISO 3696.

5.2 Petroleum ether, boiling range 40 °C to 60 °C.

5.3 Methanol.

5.4 Ammonia, 25 % aqueous solution, $\rho_{20} = 0,910$ g/ml.

5.5 Acetic acid, 100 % mass fraction, $\rho_{20} = 1,050$ g/ml.

5.6 Trisodium citrate dihydrate.

5.7 Propan-1-ol.

5.8 Ethyl acetate.

5.9 2-Methyl-2-propanol.

5.10 Propionic acid.

5.11 Eluent solution for column chromatography.

Mix 95 volumes of methanol (5.3) with 5 volumes of ammonia solution (5.4).

5.12 Acetic acid, 50 % solution in methanol.

Mix 1 volume of acetic acid (5.5) with 1 volume of methanol (5.3).

5.13 Polyamide powder, of particle size 0,05 mm to 0,16 mm.

5.14 Sand, fine granular, hydrochloric acid-washed, neutralized and calcinated.

5.15 Standard reference colours.

The purities of the standard colours may vary so it is necessary to know the purity of the colours to be used as standards. The purity shall be determined by the method AOAC 46.1.08.

NOTE Certified food colours may also be used as standards.

5.16 Standard reference solutions for thin-layer chromatography

Separately make solutions in water of each of the standard reference colours (5.15) with a standard colour content of about 1 g/l.

Prepare solutions of indigotine on the day of use. Other solutions will keep for at least 3 months (solutions of erythrosine for 1 month) when stored in the dark.

5.17 Eluent for thin-layer chromatography: solution I

Weigh, to the nearest 0,1 g, 25 g of trisodium citrate dihydrate (5.6) into a 1 000 ml one-mark volumetric flask. Dissolve in water, dilute to the mark with water and mix.

Mix 80 volumes of this citrate solution with 20 volumes of ammonia solution (5.4) and 12 volumes of methanol (5.3).

To avoid or reduce interference from safflor or saffran, it is advisable to use chromatography solution II (5.18).

5.18 Eluent for thin-layer chromatography: solution II

Mix 6 volumes of propan-1-ol (5.7) with 1 volume of ethyl acetate (5.8) and 3 volumes of water.

5.19 Eluent for thin-layer chromatography: solution III

Mix 50 volumes of 2-methyl-2-propanol (5.9) with 12 volumes of propionic acid (5.10) and 38 volumes of water.

6 Apparatus

Usual laboratory apparatus and, in particular, the following.

6.1 Mechanical or electrical homogenizing equipment, capable of homogenizing the laboratory sample.

Use a high-speed rotational cutter, or a mincer fitted with a plate with apertures not exceeding 4,0 mm in diameter.

6.2 Centrifuge tubes, of capacity 75 ml, made of glass.

6.3 Flat-bottomed flasks, of capacity 250 ml, with ground glass stoppers.

6.4 Round-bottomed flasks, of capacity 100 ml, with ground glass joint.

6.5 Centrifuge, operating at a radial acceleration of about 2 000 g_n .

6.6 Rotary evaporator.

6.7 Chromatographic column, of glass, with fritted filter and tap, of length about 20 cm, diameter about 30 mm, filter pore size 40 μm to 100 μm (porosity grade P 100 according to ISO 4793 [2]).

Put some glass wool in the column and add 1 g to 2 g of sand (5.14).

6.8 Plastics container, of volume about 10 ml, with lid.

6.9 Thin-layer plates, coated with a layer of cellulose powder of 0,10 mm thickness, or equivalent.

Ready-to-use plates are suitable.

6.10 Micropipettes, of capacity approximately 5 μl .

6.11 pH-meter, accurate to within 0,1 pH unit.

7 Sampling

Sampling is not part of the method specified in this International Standard. A recommended sampling method is given in ISO 3100-1 [1].

It is important that the laboratory receive a sample which is truly representative and has not been damaged or changed during transport or storage.

Proceed from a representative sample of at least 200 g. Store the sample in such a way that deterioration and change in composition are prevented.

8 Preparation of test sample

Homogenize the laboratory sample with the appropriate equipment (6.1). Take care that the temperature of the sample material does not rise above 25 °C. If a mincer is used, pass the sample at least twice through the equipment.

Fill a suitable airtight container with the prepared sample. Close the container and store in such a way that deterioration and change in composition of the sample are prevented. Analyse the sample as soon as practicable, but always within 24 h after homogenization.

9 Procedure

WARNINGS

If the sample contains indigotine, the temperature shall not at any time during the analysis exceed 35 °C. Indigotine partially decomposes in chromatography solution I, so chromatography solution II shall be used.

Erythrosine is sensitive to light. When pausing in the course of the analysis, solutions and plates shall be stored in the dark. The same also holds for indigotine.

9.1 Test portion

Weigh, to the nearest 0,1 g, 5 g of the prepared test sample (see clause 8) into a centrifuge tube (6.2).

For fatty samples, proceed in accordance with 9.2.

For non-fatty samples, proceed in accordance with 9.3.

9.2 Fatty samples

Add about 20 ml of petroleum ether (5.2) to the centrifuge tube and mix with a glass rod. Decant the petroleum ether.

Repeat this procedure three times.

9.3 Non-fatty samples

Add 25 ml of boiling water (see warning above) and mix. Add 25 ml of the eluent solution(5.11).

Check that the pH is $9 \pm 0,5$ using the pH-meter (6.11). If not, **adjust the pH** with acetic acid (5.5) or ammonia solution (5.4).

Mix well. Chill the sample in a freezer for 15 min (to prevent turbidity).

Centrifuge (6.5) for 10 min at a radial acceleration of about 2000 g_n .

Decant the clear solution into a flat-bottomed flask (6.3). In the case of indigotine, use a round-bottomed flask (6.4).

Add 5 ml of water to the centrifuge tube containing the residue. Mix and add 10 ml of the eluent solution (5.11). Mix and centrifuge as above.

Repeat the procedure until all colour has been extracted from the sample then combine all extracts.

Evaporate the combined extract on a water bath to about 25 ml in order to remove methanol. In the case of indigotine, use a round-bottomed flask (6.4) and the rotary evaporator (6.6) at 35 °C.

Add 25 ml of boiling water (see warnings) and mix.

9.4 Transfer of the colours to polyamide powder

Using acetic acid (5.5) or ammonia solution (5.4) adjust the pH to between 4 and 5.

Add 1 g of polyamide powder (5.13) to the warm solution (see warnings). Shake vigorously for 1 min.

Allow the powder to form a sediment.

Check that no colour remains in the solution. If the solution is coloured, add some more polyamide powder and shake vigorously.

NOTE Some natural colours (see annex B) are not entirely adsorbed on the polyamide powder, leaving the solution coloured even if all synthetic colours have been completely adsorbed. It is usually possible to decide from the type of sample whether or not such natural colours are present.

Shake and transfer the warm suspension to the chromatographic column (6.7).

Rinse the flat-bottomed flask with three 10 ml portions of hot water (see warnings) and add the rinsings, portion by portion, to the column. Wash the column another three times with 10 ml portions of hot water (see warnings) and finally three times with 5 ml of methanol (5.3). If natural colours are eluted, continue washing the column with methanol until the eluted methanol is colourless.

9.5 Elution and concentration of isolated colours

Place a flask (6.4) under the column and elute the colours from the polyamide powder with 5 ml portions of the eluent solution (5.11), at an elution volume flow rate of 2 ml/min, until the polyamide is colourless.

Evaporate the eluate to dryness using the evaporator (6.6) at a temperature of at most 35 °C (see warnings).

Add 1,0 ml or 2,0 ml of eluent solution (5.11) depending on the amount and number of colours and dissolve the residue. Transfer the colour solution to a plastics container (6.8).

9.6 Thin-layer chromatographic separation

9.6.1 Standard reference plates

Prepare three standard reference thin-layer chromatographic plates. Using a micropipette (6.10), dispense a spot of about 5 μ l (diameter < 5 mm) of each standard solution (5.16) separately on each plate (6.9). Develop these separately, one with each chromatography eluent (5.17, 5.18 and 5.19) in an unsaturated tank until the solvent front is about 10 cm to 12 cm from the starting line. Remove the plates from the tank and dry in air under a hood. Store the plates in the dark. The spots, except for that of indigotine, are stable for several years.

9.6.2 Samples

Using a micropipette (6.10), apply to a thin-layer plate (6.9) a just-visible amount of sample solution (9.5). Dry using a hair dryer. In the case of indigotine, dry in air.

Develop the plate in an unsaturated tank to a height of approximately 10 cm to 12 cm using a suitable chromatography solution (5.16, 5.17 or 5.18); i.e. the solution which gives the best separation of the colours detected in the sample (see clause 1). Sometimes it will be necessary to prepare a second sample plate and develop this in one of the other two eluents to obtain the best separation.

Remove the plate from the tank and dry in air under a hood.

Compare the sample spots with the appropriate standard reference plate (9.6.1).

It is recommended that different amounts of sample solutions be applied in the case of mixtures of colorants, because colorants may be present in various concentrations in the concentrate.

Tailing is usually caused by inadequate purification. If this is the case, adsorb the colorant again with the adsorbent, wash with hot water and remove the adsorbent as previously described.

9.7 Confirmation

Confirm the identity of the colorants by chromatographing the concentrate (9.6.2) in a mixture of standards for the colorants identified in the first chromatogram.

In case of doubt, elute the colorant from the plate with a neutral solution (water or ethanol, or 0,2 g/l ammonium acetate solution), an acid (0,1 mol/l hydrochloric acid) and an alkali (0,1 mol/l sodium hydroxide solution) and compare the absorption spectrum of the colorant to that of the standard. See the absorbance spectra shown in annex C.

10 Test report

The test report shall specify:

- all information necessary for the complete identification of the sample;
- the sampling method used, if known;
- the test method used, with reference to this International Standard;
- all operating details not specified in this International Standard, or regarded as optional, together with details of any incidents which may have influenced the test result;
- the test result obtained.

Annex A
(informative)

Synonyms and identity numbers of synthetic, water-soluble colouring agents

Table A.1

Name	Synonym 1	Synonym 2	C.I.^a	E No.^b
Tartrazine	FD&C Yellow No. 5		19140	E 102
Quinoline Yellow			47005	E 104
Sunset Yellow FCF	FD&C Yellow No. 6		15985	E 110
Amaranth	FD&C Red No. 2	Naphthol Red S	16185	E 123
Ponceau 4R	New coccine	Cochineal Red A	16255	E 124
Erythrosine	FD&C Red No. 3		45430	E 127
Patent Blue V			42051	E 131
Indigotine	FD&C Blue No. 2		73015	E 132
Brilliant Black PN			28440	E 151
Black 7984			27755	E 152
Fast Green FCF	FD&C Green No. 3		42053	c
Blue VRS			2045	c
^a C.I.: Identity number according to Colour Index [4]. ^b E No.: Current number within the European Community (EC). ^c E No. is not available.				

Annex B (normative)

Possible interference by colours

B.1 Colours which do not interfere

The following plant colours or plant extracts have been observed not to interfere with this method:

alfalfa	paprika oleoresin
annatto (bixin and norbixin)	riboflavin
anthocyanins	marigold
β -carotene	beetroot red
β -apocarotenal	mustard
β -apocarotenic acid ethyl ester	flower of tagetes
canthaxanthin	tea
chlorophyll	tomato
chlorophyllin copper complex	

B.2 Colours which may interfere

In some cases natural colours have been observed to interfere with this method. Their uses in foods and the synthetic colours of which the determination may be affected are given in Table B.1.

Table B.1

Substance	Use in foods	Interferes with analysis of
Curcumin	Spice; also used as yellow colour	Quinoline Yellow (E 104) ^a Brilliant Black PN (E 151) ^a Black 7984 (E 152) ^a
Saffran	Spice (too expensive for use as a colour)	Erythrosine (E 127) ^b Quinoline Yellow (E 104) ^{a, b} Brilliant Black PN (E 151) ^{a, b} Black 7984 (E 152) ^b
Safflor	Substitute for saffran	Tartrazine (E 102) ^b
^a The interferences are minor and may be considered negligible.		
^b To avoid or reduce interference from safflor or saffran, it is advisable to use chromatography solution II (5.18).		

Annex C (informative)

Absorbance spectra

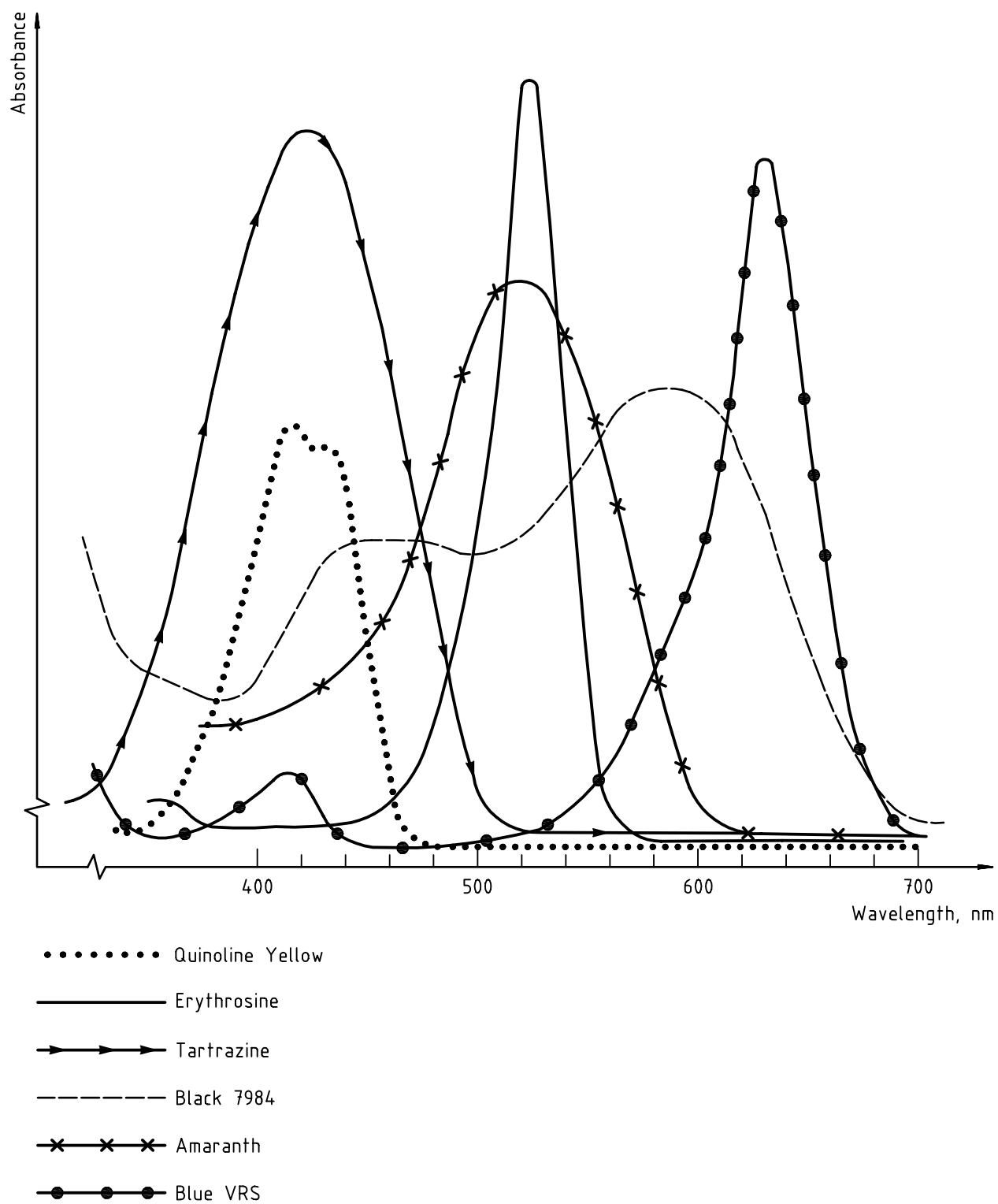


Figure C.1 — Absorbance spectra of Amaranth, Black 7984, Blue VRS, Erythrosine, Tartrazine and Quinoline Yellow

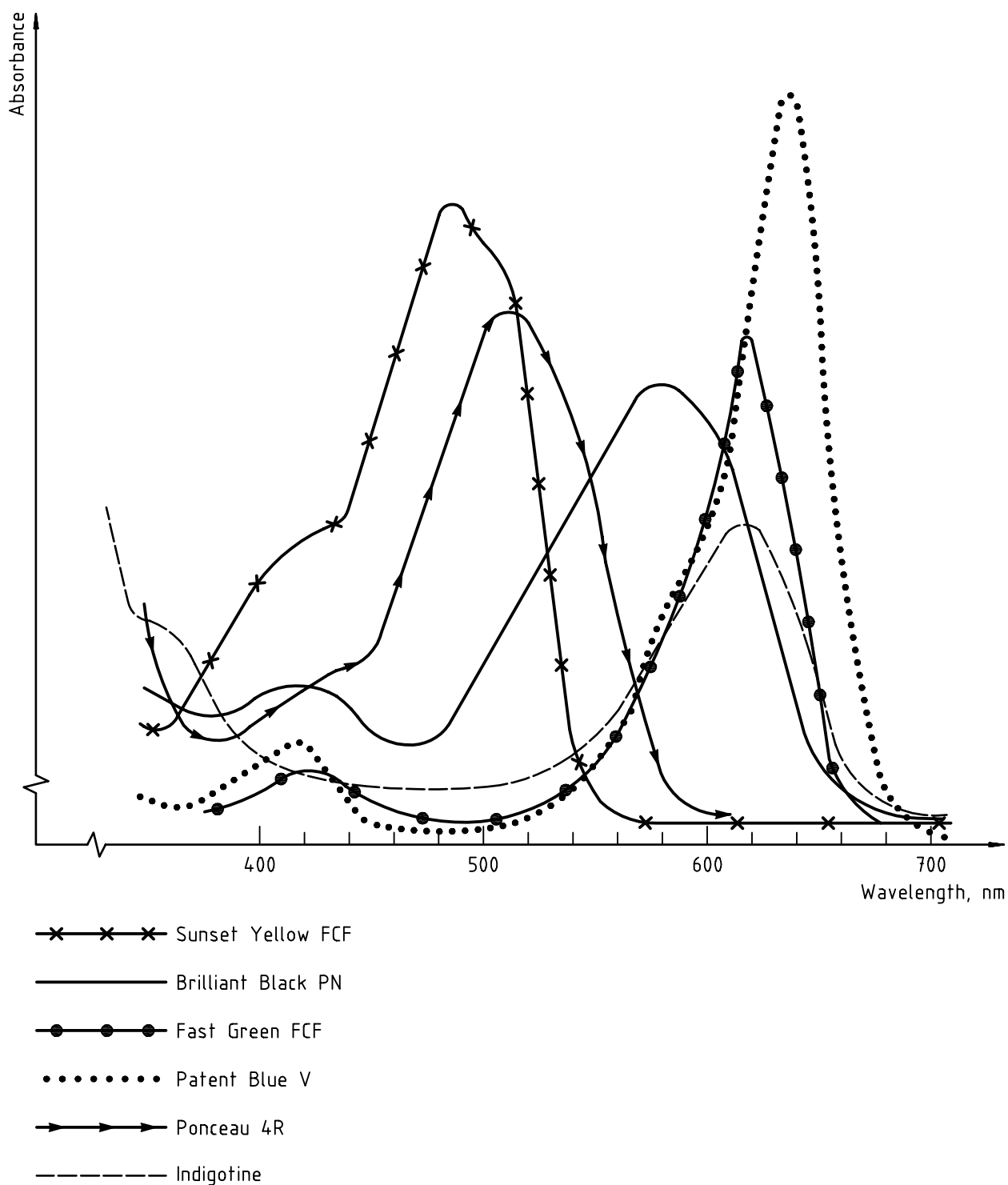


Figure C.2 — Absorbance spectra of Brilliant Black PN, Fast Green FCF, Indigotine, Patent Blue V, Ponceau 4R and Sunset Yellow FCF

Bibliography

- [1] ISO 3100-1, *Meat and meat products — Sampling and preparation of test samples — Part 1: Sampling.*
- [2] ISO 4793, *Laboratory sintered (fritted) filters — Porosity grading, classification and designation.*
- [3] *Colours, synthetic, water-soluble. Semi-quantitative determination by chromatography and spectrophotometry.* Nordic Committee on Food Analysis (NMKL), No. 134, 1990.
- [4] *Colour Index*, 3rd edition, 1971.

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Review of Indian Standards

Amendments are issued to standards as the need arises on the basis of comments. Standards are also reviewed periodically; a standard along with amendments is reaffirmed when such review indicates that no changes are needed; if the review indicates that changes are needed, it is taken up for revision. Users of Indian Standards should ascertain that they are in possession of the latest amendments or edition by referring to the latest issue of 'BIS Catalogue' and 'Standards: Monthly Additions'.

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